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GENETIC DIVERSITY ANALYSIS OF BRINJAL (*SOLANUM MELONGENA* L.) USING RAPD MARKERS

Shetul Muna Parvin
Post graduate student
Department of Horticulture
Bangladesh Agricultural University
Mymensingh-2202, Bangladesh

Md. Golam Rabbani
Professor
Department of Horticulture
Bangladesh Agricultural University
Mymensingh-2202, Bangladesh

Md. Abu Zafur Al Munsur
PhD Fellow, Department of
Agronomy, Sher-e-Bangla
Agricultural University, Dhaka,
Bangladesh.
Additional Deputy Director (LR)
Department of Agricultural
Extension, Khamarbari, Dhaka,
Bangladesh.

Abstract— An experiment was conducted at the Horticulture Farm, Department of Horticulture, Bangladesh Agricultural University, Mymensingh to study the variability of 12 selected accessions of brinjal using morphological traits and RAPD. Molecular characterization and diversity study was conducted using RAPD markers. Amplification with 5 decamer primers produced 54 RAPD markers of which 49 (89.56%) were polymorphic and 5 (10.44%) were monomorphic. The highest genetic distance (0.9445) and the lowest genetic identity (0.3889) were observed in the accessions SM59 vs. SM11 combination. The lowest genetic distance (0.1388) was observed between the accessions SM22 and SM29 and higher genetic identity was also observed in this combination. Dendrogram was constructed from genetic distance and 12 brinjal accessions fall in five clusters. Thus the RAPD analysis provided an effective method in investigating genetic variation and relatedness in twelve brinjal accessions.

Keywords— Brinjal, Genetic diversity, RAPD, *Solanum melongena*,

I. INTRODUCTION

Brinjal (*Solanum melongena* L.) is an important solanaceous vegetable crop worldwide and also in Bangladesh. It has high nutritional value in respect of calorie, iron, phosphorus and riboflavin contents than tomato [1] (Shaha, 1989). It is the second most important vegetable crop after potato in Bangladesh in respect of total area, and third in respect of production [2]. The fruits, leaves, stem and roots of brinjal are widely used in ayurvedic medicine for the patients suffering from diabetes, asthma, tooth ache, cholera, bronchitis and

dysentery. Brinjal has potentiality as raw material in pickle making and in dehydration industries [3] (Singh *et al.*, 1963). Due to its high nutritional value, taste, easy cooking quality and year round availability, brinjal has become a widely consumed and popular vegetable in Bangladesh.

Although it is the major native vegetables in our country, very little attention has been given for its improvement and conservation. Among the cultivated varieties, wide range of genetic variability exists which would disappear in course of time. Therefore, those germplasm should be properly evaluated and conserved for future use. Variability is the touch-stone to a breeder to evolve high yielding varieties through selection either from the existing accessions or from the segregates of different crosses. Hence, the genetic information on yield and yield contributing characters need to be properly assessed for its improvement. Also, similarity or duplication of the germplasm should be avoided for mid or long term conservation of the collected germplasm. For rapid and efficient plant breeding program, knowledge of interrelationship among yield and yield contributing characters is also necessary.

Genetic diversity studies based on morphological traits and molecular marker studies enable the evaluation of genetic variation prior to the initiating of a new breeding programme. Molecular markers have appeared to be very useful tools for assessing bio-diversity and revealing differences and relationship between taxa [4] (Dore, 2001). Crop improvement programme based on phenotypic markers are generally slower than desired, because phenotypic assessment does not always reflect true genotypic performance. Moreover phenotypic markers are affected by environment and may



have weak linkage with the target trait(s). Molecular markers like SSR, RFLP, RAPD and AFLP. are being developed and applied to gene banks and plant breeding programs, particularly for the identification of accessions, the detection of genetic relationships and introduction of those markers based on polymerase chain reaction (PCR) technology has led to the development of several novel genetic assays that can be used for many purposes in plant genetic analysis such as cultivar identification, gene mapping and so on. The study was taken to characterize the brinjal germplasm at molecular level through RAPD markers.

II. MATERIALS AND METHODS

The experiment was conducted at Horticulture Farm, Vegetable and Fruit Germplasm Laboratory, Department of Horticulture and Central Laboratory, Bangladesh Agricultural University, Mymensingh.

Planting materials used for the experiment

Twelve brinjal accessions were included in this study for genetic diversity analysis using morphological traits and molecular markers that were collected from different parts of Bangladesh.

Plant genomic DNA isolation

Genomic samples were extracted from vigorous, young actively growing leaf tissue following modified [5] method. Approximately 0.1g of leaf tissues was cut into small pieces and digested by 800 μ l extraction buffer. The ground sample was vortexed for 20 seconds (IUCHI Automatic Lobo mixture, Japan) and incubated at 65°C for 5 min in hot water bath (JP Selecta^(R)). The extract was centrifuged for 10 min at 14000 rpm with a micro centrifuge to allow precipitation of the cell debris. The upper aqueous phase of about 600 μ l was transferred to another eppendorf tube. For purification, equal volume 600 μ l of Phenol: Chloroform: Isoamyl alcohol (v: v: v=25: 24: 1) was added to the tube and vortexed for few seconds. The upper aqueous phase was recovered and DNA was precipitated with 800 μ l of absolute ethanol (100%). At this point, DNA became visible as white strands by flicking the tube several times with fingers. DNA was pelleted by centrifugation for 3 min at 14000 rpm. Then the liquid was discarded completely and re-precipitation of the DNA solution was done by adding 400 μ l of 70% ethanol with 20 μ l 3 M sodium acetate and pelleted by centrifugation for 3 min at 14000 rpm. The liquid was removed completely by pouring and blotting the open tube end on fresh tissue paper. The pellets were then air dried and dissolved in an appropriate volume (20 μ l-50 μ l) of TE buffer, pH 8.0. Finally, the DNA samples were stored at -20°C. Isolated DNA prepared by above protocol often contains a large amount of RNA and pigments that can usually cause spuriously high estimation of DNA concentration on a spectrophotometer. For this 1.8% agarose gel was used for assessing both the quantity of the genomic DNA.

Selection of primers

A final subset of five primers out of total of twenty decamer primers (Operon Technologies, Inc., Alameda, California,

USA) exhibiting better quality banding patterns was selected for the analysis of whole sample set of the 12 accessions of brinjal (Table 1). To confirm the reproducibility of RAPD markers, the selected five primers were screened three times on the same samples. It is best to do a series of RAPD reactions using a couple of primers and a set of serial dilutions of each genomic DNA identify empirically the useful range of DNA concentrations, for which reproducible RAPD patterns were obtained.

Table 1. Parameters of the operon random primers used for brinjal accessions

Primer code	Sequences (5'-3')	(G +C) %
OPA-04	AATCGGGCTG	60
OPC-05	GATGACCGCC	70
OPC-06	GAACGGACTA	60
OPL-05	ACGCAGGCAC	60
OPL-07	AGGCGGGAAC	70

PCR amplification for RAPD

The amplification of PCR conditions was based on with some modifications [6]. PCR reactions were performed on each DNA sample in a 10 μ l reaction mix containing 1 μ l of 10 \times Ampli Taq polymerase buffer, 0.25 μ l of 10 μ M primer, 1 μ l of 250 μ M dNTPs, 1 unit of Ampli Taq DNA polymerase (Bangalore Genei, India) and 4 μ l (100 ng) of genomic DNA and suitable amount of sterile deionized water. The PCR was performed in an oil-free thermal cycler (Master Cycler Gradient, Eppendorf). The reaction mix was preheated at 94°C for 3 minutes followed by 40 cycles consisted of 1 min denaturation at 94°C, 1 min at 34°C for annealing and 2 min at 72°C for elongation or extension. After the last cycle, a final step of 7 min at 72°C was added to allow complete extension of all amplified fragments of DNA.

The amplified product from each sample was separated electrophoretically on 1.8% agarose gel containing 6- μ l ethidium bromide in 1 X TBE buffer at 100V for 60 minutes. Loading dye (2.0 μ l) was added to the PCR products and loaded in the wells. Two molecular weight marker PUC and 100 bp DNA ladder were also loaded on either side of the gel. Under Ultra Violet (UV) light on a transilluminator RAPD bands were observed and documented by taking photograph using a Gel Cam Polaroid camera.

RAPD data analysis: All distinct bands or fragments (RAPD markers) were thereby given identification numbers according to their position on gel and scored visually on the basis of their presence (1) or absence (0), separately for each individual and each primer. The scores obtained using all primers in the RAPD analysis were then pooled to create -a single data matrix. This was used to estimate polymorphic loci [7] (Nei's, 1973), gene diversity, population differentiation (G_{st}), gene flow (N_m), genetic distance (D) and to construct a UPGMA dendrogram among populations using a computer program, POPGENE (Version 1.31) [8] (Yeh *et al.*, 1999). The similarity index values (SI) between the RAPD profiles of any two individuals on the same gel were calculated from RAPD markers according to the this formula: Similarity index (SI) = $2 N_{xy}/N_x + N_y$ Where, N_{xy} is the total number of RAPD bands

shared by individuals x and y respectively, and N_x and N_y are the number of bands in individual x and y , respectively [9] (Chapco *et al.*, 1992). The SI value range from 0 to 1. When $SI=1.0$, the two DNA profiles are identical and when SI is 0.0, there are no common bands between the two profiles. Within population similarity (S_i) was calculated as the average of SI across all possible comparisons between individuals within a population. Between population similarity (S_{ij}) was calculated as the average similarity between each paired individuals of population i and j [10] (Lynch, 1991).

III. EXPERIMENT AND RESULT

Primer selection and RAPD pattern

Among the 20 primers initially tested, five decamer primers (OPA-4, OPC-5, OPC-6, OPL-5, and OPL-7) were used in RAPD analysis of twelve brinjal accessions. These primers yielded comparatively maximum number of amplification products with high intensity with minimal smearing, good resolution and also clear bands. The number of fragments amplified per primer varied. The five primers generated 54 distinct bands with size ranging from 140-1500 bp, of which 49 (89.56%) were considered as polymorphic and 5 bands (10.44%) were monomorphic amongst the 12 brinjal accessions. Among the 5 primers, OPC-5 and OPC-6 produced maximum number of polymorphic bands (12) that indicated a high level of polymorphism and other primers OPA-4, OPL-7 and OPL-5 produced 10, 8 and 7 respectively (Table 2).

Table 2: RAPD primers with corresponding bands score and their size range together with polymorphic bands observed in 12 brinjal accessions

Primer codes	Total number of bands scored	Size ranges (bp)	Number of polymorphic bands	Proportion of polymorphic loci (%)
OPA-04	11	350-1400	10	90.00
OPC-05	13	140-1500	12	90.31
OPC-06	12	250-1440	12	100
OPL-05	8	250-1500	7	87.50
OPL-07	10	175-1440	8	80
Total	54		49	447.81
Average	10.8		9.8	89.562

The level of polymorphism (89.56%) indicates the effectiveness of RAPD technique to detect substantial amount of polymorphism or diversity among different accessions of brinjal. The banding patterns of 12 brinjal accessions using primers OPA-4, OPC-5, OPC-6, OPL-7 and OPL-7 are shown in plate 2-6.

Plate 1. RAPD profiles of 12 brinjal accessions using primer OPC-06. M: Molecular weight marker (100 bp DNA ladder in left side and pUC in right side).

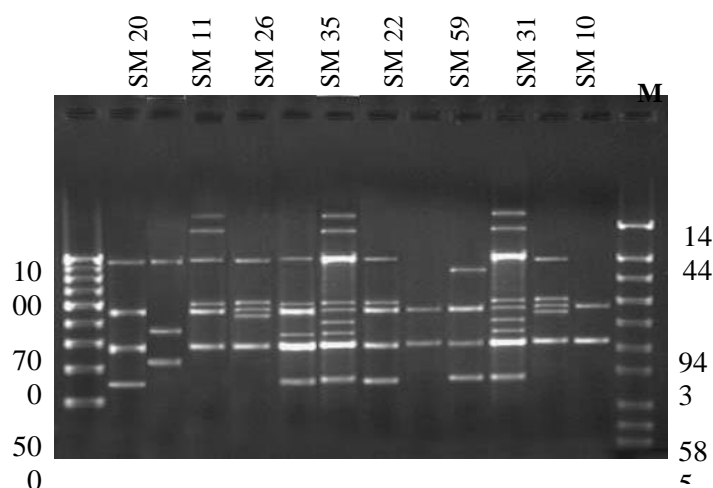
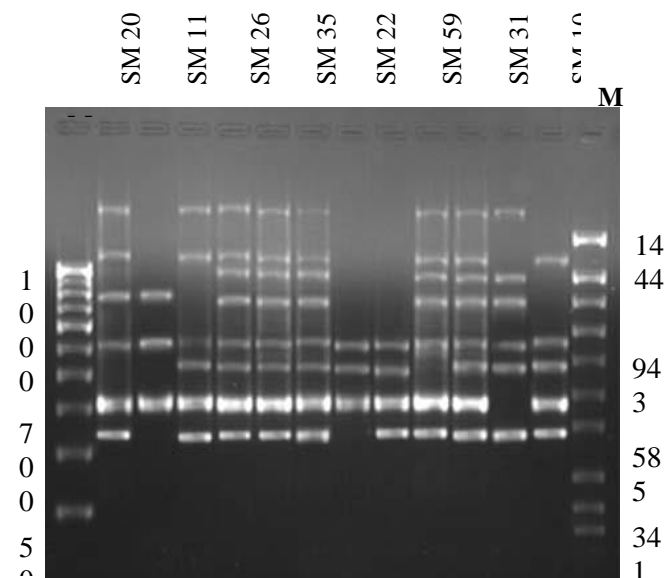


Plate 2. RAPD profiles of 12 brinjal accessions using primer OPL-05. M: Molecular weight marker (100 bp DNA ladder in left side and pUC in right side).



Frequency of polymorphic loci

The DNA polymorphisms were detected according to the presence and absence of band. Absence of band may be caused by failure of primer to anneal a site in some individuals due to nucleotide sequences difference or by insertions or deletions between primer sites [11] (Clark and Lanigan, 1993). Frequencies of maximum number of polymorphic loci were found to be high with the exception of OPC06-4 (0.0833) and no frequencies were found in OPA04-7 (0.000) and OPL05-5 (0.0000) (Table 3).



Table 3: Frequencies of polymorphic RAPD markers in 12 brinjal accessions

RAPD Markers	Gene frequency	RAPD Markers	Gene frequency
OPA04-1	0.1667	OPC06-4	0.0833
OPA04-2	0.5000	OPC06-5	0.5833
OPA04-3	0.7500	OPC06-6	0.9167
OPA04-4	0.8333	OPC06-7	0.1667
OPA04-5	0.5000	OPC06-8	0.1667
OPA04-6	0.5000	OPC06-9	0.3333
OPA04-7	0.0000	OPC06-10	0.9167
OPA04-8	0.2500	OPC06-11	0.0833
OPA04-9	0.5833	OPC06-12	0.5000
OPA04-10	0.6667	OPL05-1	0.6667
OPA04-11	0.2500	OPL05-2	0.6667
OPC05-1	0.5833	OPL05-3	0.5000
OPC05-2	0.2500	OPL05-4	0.6667
OPC05-3	0.5833	OPL05-5	0.0000
OPC05-4	0.5000	OPL05-6	0.7500
OPC05-5	0.6667	OPL05-7	0.9167
OPC05-6	0.1667	OPL05-8	0.8333
OPC05-7	0.5833	OPL07-1	0.5833
OPC05-8	0.0000	OPL07-2	0.5833
OPC05-9	0.6667	OPL07-3	0.9167
OPC05-10	0.6667	OPL07-4	0.1667
OPC05-11	0.5000	OPL07-5	0.9167
OPC05-12	0.4167	OPL07-6	0.4167
OPC05-13	0.8333	OPL07-7	0.5000
OPC06-1	0.2500	OPL07-8	0.0000
OPC06-2	0.2500	OPL07-9	0.0000
OPC06-3	0.7500	OPL07-10	0.3333

Genetic variation

The estimate of [7] Nei's (1973) genetic diversity for entire accessions of brinjal was 0.36 and Shannon's information index was 0.54. There was a high level of genetic variation among the studied accessions of brinjal from the proportion of polymorphic loci point of view (Table 4).

Table 4: Summary of genetic diversity statistics for all loci in 12 brinjal accessions

Loci	Gene diversity (h)	Shanon information index (i)	Loci	Gene diversity (h)	Shanon information index (i)
OPA04-1	0.2778	0.4506	OPC06-4	0.1528	0.2868
OPA04-2	0.5000	0.6931	OPC06-5	0.4861	0.6792
OPA04-3	0.3750	0.5623	OPC06-6	0.1528	0.2868
OPA04-4	0.2778	0.4506	OPC06-7	0.2778	0.4506
OPA04-5	0.5000	0.6931	OPC06-8	0.2778	0.4506
OPA04-6	0.5000	0.6931	OPC06-9	0.4444	0.6365
OPA04-7	0.0000	0.0000	OPC06-10	0.1528	0.2868
OPA04-8	0.3750	0.5623	OPC06-11	0.1528	0.2868
OPA04-9	0.4861	0.6792	OPC06-12	0.5000	0.6931
OPA04-10	0.4444	0.6365	OPL05-1	0.4444	0.6365
OPA04-11	0.3750	0.5623	OPL05-2	0.4444	0.6365
OPC05-1	0.4861	0.6792	OPL05-3	0.5000	0.6931
OPC05-2	0.3750	0.5623	OPL05-4	0.4444	0.6365
OPC05-3	0.4861	0.6792	OPL05-5	0.0000	0.0000
OPC05-4	0.5000	0.6931	OPL05-6	0.3750	0.5623
OPC05-5	0.4444	0.6365	OPL05-7	0.1528	0.2868
OPC05-6	0.2778	0.4506	OPL05-8	0.2778	0.4506
OPC05-7	0.4861	0.6792	OPL07-1	0.4861	0.6792
OPC05-8	0.0000	0.0000	OPL07-2	0.4861	0.6792
OPC05-9	0.4444	0.6365	OPL07-3	0.1528	0.2868
OPC05-10	0.4444	0.6365	OPL07-4	0.2778	0.4506
OPC05-11	0.5000	0.6931	OPL07-5	0.1528	0.2868

OPC05-12	0.4861	0.6792	OPL07-6	0.4861	0.6792
OPC05-13	0.2778	0.4506	OPL07-7	0.5000	0.6931
OPC06-1	0.3750	0.5623	OPL07-8	0.0000	0.0000
OPC06-2	0.3750	0.5623	OPL07-9	0.0000	0.0000
OPC06-3	0.3750	0.5623	OPL07-10	0.4444	0.6365
Mean				0.3457	0.5092
Standard deviation				0.1600	0.2114

Genetic distance and genetic identity

Pair-wise comparisons of [12] Nei's (1972) genetic distance (GD) between brinjal accessions were computed from combined data for the five primers and the values ranged from 0.1388 to 0.9445 (Table 6). Comparatively higher genetic distance was observed between SM59 vs. SM11 (0.9445) while the lowest genetic distance 0.1388 was observed between the accessions SM 22 and SM29. SM35 vs. SM11 and SM44 vs. SM11 (0.8535) accession pairs than other accession combinations. Considering the genetic distance values the results indicate that some accessions of brinjal are genetically different from each other and some are tend to be similar. Genetic identity between varieties was found for the 5 primers, ranged from 0.3889 to 0.8704. The highest genetic identity (0.8704) was found in SM22 vs. SM29. The lowest (0.3889) was observed between SM11 vs. SM59 (Table 5).

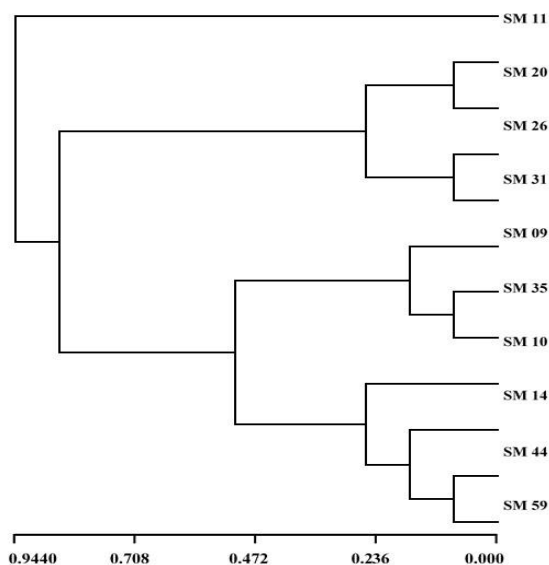
Table 5. Summary of [12] genetic identity (above diagonal) and genetic distance (below diagonal) values between 12 brinjal germplasm

Genotypes	SM 20	SM 11	SM 26	SM 35	SM 22	SM 59	SM 31	SM 10	SM 44	SM 29	SM 09	SM 14
SM 20	****	0.592	0.851	0.611	0.648	0.537	0.703	0.592	0.648	0.5926	0.722	0.611
SM 11	0.523	***	0.518	0.425	0.500	0.388	0.481	0.481	0.425	0.4444	0.500	0.537
SM 26	0.160	0.656	****	0.611	0.611	0.537	0.666	0.555	0.537	0.5926	0.685	0.611
SM 35	0.492	0.853	0.492	***	0.740	0.703	0.648	0.759	0.703	0.7222	0.666	0.703
SM 22	0.433	0.693	0.492	0.300	****	0.777	0.574	0.648	0.851	0.8704	0.518	0.629
SM 59	0.621	0.944	0.621	0.351	0.251	****	0.574	0.648	0.703	0.8333	0.518	0.629
SM 31	0.351	0.730	0.405	0.433	0.555	0.555	****	0.740	0.537	0.5185	0.722	0.611
SM 10	0.523	0.730	0.587	0.275	0.433	0.433	0.300	****	0.685	0.6296	0.611	0.833
SM 44	0.433	0.853	0.621	0.351	0.160	0.351	0.621	0.378	****	0.7593	0.444	0.666
SM 29	0.523	0.810	0.523	0.325	0.138	0.182	0.656	0.462	0.275	****	0.537	0.611
SM 09	0.325	0.693	0.378	0.405	0.656	0.656	0.325	0.492	0.810	0.6217	***	0.555
SM 14	0.492	0.621	0.492	0.351	0.462	0.462	0.492	0.182	0.405	0.4925	0.587	****

UPGMA dendrogram

The Unweighted Pair Group Method of Arithmetic Means (UPGMA) dendrogram based on [12] genetic distance was constructed. This measurement indicated the segregation of the 12 accessions of brinjal into 2 main clusters. SM 11 alone formed separate cluster 'A'. Rest 11-brinjal germplasm grouped in cluster 'B' (Fig.1). SM11 belonged to the cluster A. In cluster B, SM20, SM26, SM31 and SM09 formed sub cluster B-I while SM35, SM10, SM14, SM44, SM59, SM22 and SM29 formed sub cluster B-II. Further, the varieties of sub cluster B-II were divided into sub sub-cluster 3 and subsub-cluster 4. SM35, SM 10 and SM 14 formed sub sub-cluster 3 and SM44, SM59, SM22 and SM29 belonged to sub sub-cluster 4.

Fig.1. Unweighted pair group method of arithmetic mean (UPGMA) dendrogram based on [12] genetic distance, summarizing data on differentiation in 12 brinjal accessions according to RAPD analysis



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IV. CONCLUSION

RAPD markers produced 89.56% polymorphic band and the highest genetic distance (0.9445) were observed in the accessions SM59 vs SM11 combination. So, It can be concluded that this cultivar used for further variety development.

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